

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
21 February 2002 (21.02.2002)

PCT

(10) International Publication Number
WO 02/13766 A2

(51) International Patent Classification⁷: A61K [US/IL]; 3 Ha'absor, Apt. #2, Herzleya (IL). BIALER, Meir [IL/IL]; 36 Hehalutz Street, 96222 Jerusalem (IL).

(21) International Application Number: PCT/US01/25919

(22) International Filing Date: 17 August 2001 (17.08.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/225,973 17 August 2000 (17.08.2000) US
60/225,977 17 August 2000 (17.08.2000) US

(71) Applicants (for all designated States except BB, US): TEVA PHARMACEUTICAL INDUSTRIES, LTD. [IL/IL]; 5 Basel Street, P.O. Box 3190, 49131 Petah Tiqva (IL). YISSUM RESEARCH DEVELOPMENT COMPANY OF THE HEBREW UNIVERSITY OF JERUSALEM [IL/IL]; 36 Jabotinsky Street, 91042 Jerusalem (IL).

(71) Applicant (for BB only): TEVA PHARMACEUTICALS USA, INC. [US/US]; 1090 Horsham Road, North Wales, PA 19454 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): SHIRVAN, Mitchell

(74) Agent: WHITE, John, P.; Cooper & Dunham LLP, 1185 Avenue of the Americas, New York, NY 10036 (US).

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (regional): ARJPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— without international search report and to be republished upon receipt of that report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

WO 02/13766 A2

(54) Title: USE OF DERIVATIVES OF VALPROIC ACID AMIDES AND 2-VALPROENIC ACID AMIDES FOR THE TREATMENT OR PREVENTION OF PAIN AND/OR HEADACHE DISORDERS

(57) Abstract: A method for the treatment or prevention of pain and/or a headache disorder using a derivative of a valproic acid amide or a 2-valproenic acid amide, as well as pharmaceutical compositions comprising these derivatives or compounds.

Applicants: Daniella Licht et al.
Serial No.: 10/773,442
Filed: February 5, 2004
Exhibit 4

USE OF DERIVATIVES OF VALPROIC ACID AMIDES
AND 2-VALPROENIC ACID AMIDES FOR THE TREATMENT
OR PREVENTION OF PAIN AND/OR HEADACHE DISORDERS

5

This application claims the benefit of U.S. Provisional Application No. 60/225,973, filed August 17, 2000 and U.S. Provisional Application No. 60/225,977, filed August 17, 2000.

10

Throughout this application, various references are referenced by short citations within parenthesis. Full citations for these references may be found at the end of the specification, immediately preceding the claims. These references, in their entireties, are hereby incorporated by reference to more fully describe the state of the art to which this invention pertains.

15

Field of the Invention

20

Disclosed is a method for the treatment or prevention of pain and/or headache disorders, such as migraines, using derivatives of valproic acid amides and 2-valproenic acid amides.

Background of the Invention

25

Pain is considered to play a basic physiological role in the detection and localization of tissue damage or potentially damaging physiological processes. Pain has been broadly classified as somatogenic, where a physiological explanation can be found, or psychogenic, where the physiological explanation is not known (The Merck Manual of Diagnosis and Therapy).

30

One example of a somatogenic pain is neuropathic pain. Generally, neuropathic pain is described as a pain which results from a dysfunction in the central or peripheral nervous system (Tremont-Lukats, I. et al.; Woolf, C. and Mannion, R.).

35

-2-

The pain can be both chronic and acute, and can also be evoked by noxious stimuli, also referred to as hyperalgesia, or by non-noxious stimuli referred to as allodynia (Attal, N.). Allodynia and hyperalgesia can have mechanical causes (dynamic or static), or a thermal cause. Examples of neuropathic pain include: all the painful peripheral neuropathies and specifically diabetic peripheral neuropathy; postherpetic neuralgia; and trigeminal neuralgia. Trigeminal neuralgia, for example, is the most common neuralgic syndrome in the elderly. The initial drug of choice is carbamazepine. For other types of pain, such as postherpetic neuralgia and painful diabetic neuropathy, amitriptyline is most commonly used. Other types of somatogenic pain that may have neuropathic components include cancer pain, postoperative pain, low back pain, complex regional pain syndrome, phantom pain, HIV pain, arthritis (osteo-arthritis and rheumatoid arthritis) pain and migraines.

Pain may also be a symptom of headache disorders. Migraines constitute one of the four major categories of primary headaches (International Headache Society; Silberstein, S.D. et al.). The other three types of primary headaches are tension-type, cluster and a miscellaneous-type (International Headache Society; Silberstein, S.D. et al.). One current view is that there is a continuous spectrum of headache severity ranging from mild tension headaches to severe migraines. Others consider tension headaches and migraines to be distinct entities.

Migraines are considered to be a familial disorder characterized by periodic pulsatile headaches. (Principles of Neurology). Migraines are found in about 4% of the male population and 7% of the female population. Migraines can occur in the presence or absence of an aura. An aura is a complex of focal neurological symptoms which may precede or

-3-

accompany a migraine attack (Silberstein, S.D. et al.). Auras can be characterized by visual, sensory, or motor phenomenon, and may also involve language or brainstem disturbances (Silberstein, S.D. et al.).

5 A major theory regarding the pain of migraines is that it stems from a form of sterile neurogenic inflammation (Moskowitz, M.A. and Cutrer, F.M.). The neurogenic inflammation results in the leakage of plasma proteins into the dura mater, which can be quantified by measuring the leakage of radioactive albumin
10 (Suzzi, M.C. and Moskowitz, M.A.).

Drugs used in the treatment of headache disorders such as migraines originate from a broad range of different drug categories. These include: 5-hydroxytryptamine agonists (5-HT₁ agonists); dihydroergotamine; ergotamine; anti-emetics; 15 anxiolytics; non-steroidal anti-inflammatory drugs; steroids; major tranquilizers; narcotics; beta-blockers; calcium channel blockers; anti-depressants; and anti-epileptic drugs. However, not all of the drugs in these categories are truly effective.
20 Considering all of the drugs which are effective, there is still a need for more efficacious drugs, as well as anti-migraine treatments with less side effects.

U.S. Patent 5,585,358 describes a series of derivatives of 25 valproic acid amides and 2-valproenic acid amides for the treatment of epilepsy and other neurological disorders. However, U.S. Patent 5,585,358 does not teach or suggest the use of derivatives of valproic acid amides and 2-valproenic acid amides for the treatment or prevention of pain or headache
30 disorders.

-4-

Summary of the Invention

The subject invention provides a method of treating or preventing pain and/or a headache disorder in a subject comprising the administration of a therapeutically effective amount of a derivative of a valproic acid amide or a 2-valproenic acid amide, to thereby treat or prevent the pain and/or headache disorder. In addition, the subject invention contains pharmaceutical compositions comprising these derivatives.

10

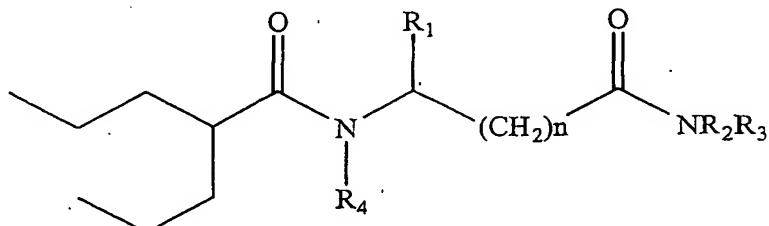
Description of the Drawings

Figure 1 presents the effects of the administration of VGD (valproylglycine amide or Compound 1) versus MC (methyl cellulose or vehicle) in the Chung model of neuropathic pain.

15

Detailed Description of the Invention

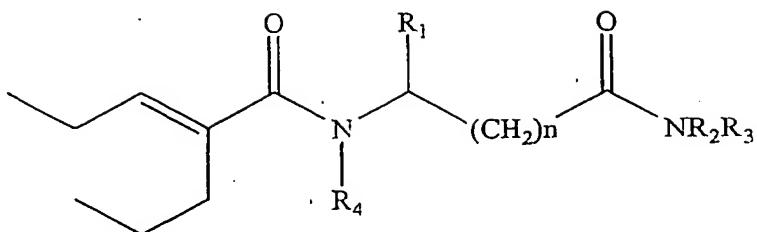
The subject invention provides a method of treating subject suffering from pain comprising periodically administering to the subject a therapeutically effective amount of a compound having the following structure:



30

or

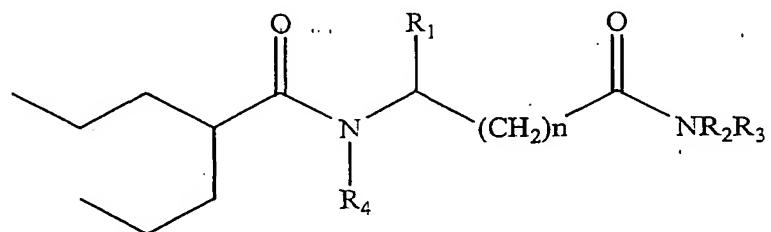
-5-



5

wherein R₁, R₂, R₃ and R₄ are independently the same or different and are hydrogen, a linear or branched C₁-C₆ alkyl group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3, so as to thereby treat the subject's pain.

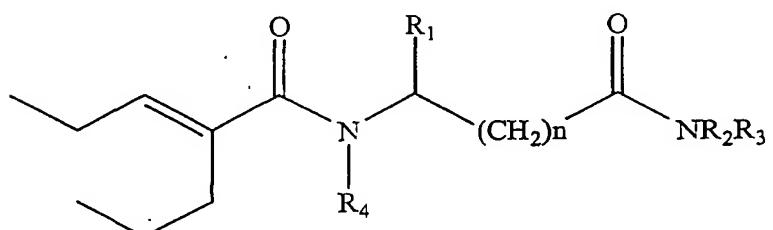
The subject invention also provides a method of preventing pain in a subject predisposed to suffering from pain comprising periodically administering to the subject a prophylactically effective dose of a compound having the following structure:



20

or

25



30

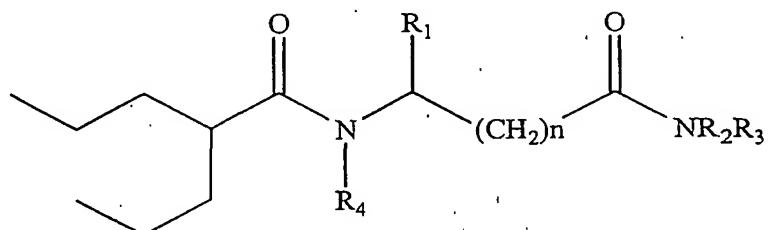
wherein R₁, R₂, R₃ and R₄ are independently the same or different and are hydrogen, a linear or branched C₁-C₆ alkyl

-6-

group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3, so as to thereby prevent pain in the subject.

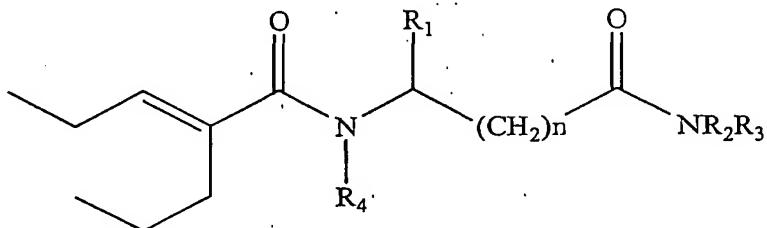
5 In addition, the subject invention provides a method of treating a subject suffering from a headache disorder comprising periodically administering to the subject a therapeutically effective dose of a compound having the following structure:

10



or

20



25

wherein R₁, R₂, R₃ and R₄ are independently the same or different and are hydrogen, a linear or branched C₁-C₆ alkyl group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3, so as to thereby treat the headache disorder.

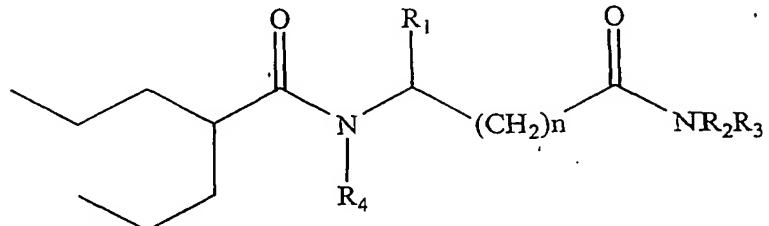
30

The subject invention further provides a method of preventing a headache disorder in a subject predisposed to suffering from a headache disorder comprising periodically administering to

-7-

the subject a prophylactically effective dose of a compound having the following structure:

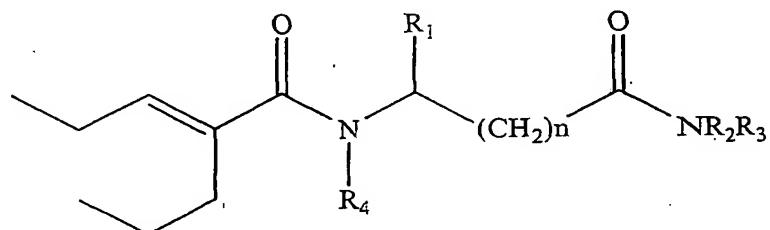
5



or

10

15

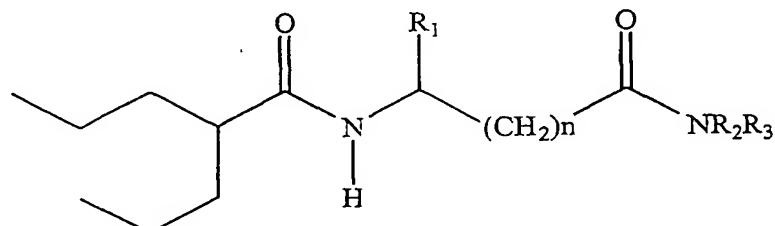


wherein R_1 , R_2 , R_3 and R_4 are independently the same or different and are hydrogen, a linear or branched C_1-C_6 alkyl group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3, so as to thereby prevent the headache disorder in the subject.

25

In one embodiment, the compound has the following structure:

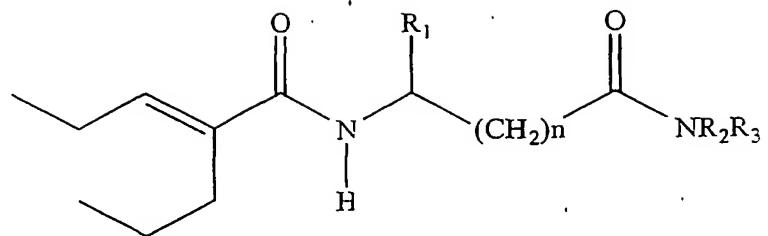
30



or

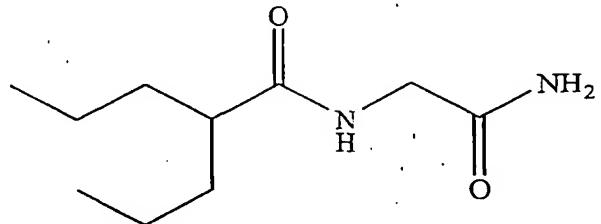
-8-

5



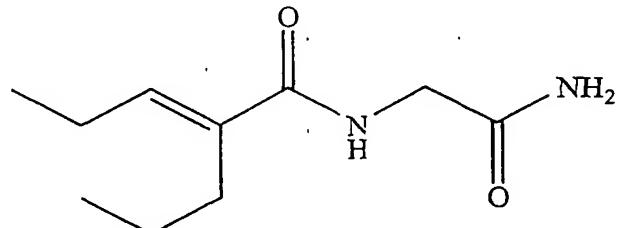
10 In an additional embodiment, the compound is N-(2-n-propylpentanoyl)-glycinamide and has the structure:

15



20 In another embodiment, the compound is N-(2-n-propylpent-2-enoyl)-glycinamide and has the structure:

25



30

In one embodiment, the pain is acute. In another embodiment, the pain is chronic. In a further embodiment, the pain is somatogenic pain. In a preferred embodiment, the pain is neuropathic pain.

-9-

The headache disorder may be a migraine.

The headache disorder may be a cluster headache.

The headache disorder may be a tension-type headache.

5

The headache disorder may be a miscellaneous-type headache.

The subject may be a human being.

10 In one embodiment, one or more of R_1 , R_2 , R_3 , or R_4 is a linear chain C_1 - C_6 alkyl group. In another embodiment, one or more of R_1 , R_2 , R_3 , or R_4 is a branched chain C_1 - C_6 alkyl group. In yet another embodiment, one or more of R_1 , R_2 , R_3 , or R_4 is a benzyl, alkylbenzyl, hydroxybenzyl, alkoxycarbonylbenzyl, 15 aryloxycarbonylbenzyl, carboxybenzyl, nitrobenzyl, cyanobenzyl, or halobenzyl group. In still another embodiment, one or more of R_1 , R_2 , R_3 , or R_4 is a phenyl, naphthyl, anthracenyl, pyridinyl, indolyl, furanyl, alkylphenyl, hydroxyphenyl, alkoxycarbonylphenyl, aryloxycarbonylphenyl, nitrophenyl, 20 cyanophenyl, halophenyl group, mercaptophenyl, or aminophenyl group.

25 The subject invention also provides a method of treating a subject suffering from neuropathic pain comprising administering to the subject 500 mg of N-(2-n-propylpentanoyl)glycinamide six times per day so as to thereby treat the subject's neuropathic pain.

30 In addition, the subject invention provides a method of preventing neuropathic pain in a subject predisposed to suffering from neuropathic pain comprising administering to the subject 500 mg of N-(2-n-propylpentanoyl)glycinamide six times

-10-

per day so as to thereby prevent neuropathic pain in the subject.

Some of the compounds used in this invention possess chiral centers. It is a further embodiment of this invention that these compounds may comprise substantially pure D or L enantiomers or racemic mixtures. It is to be understood that compounds of this invention may be of the E-(trans) or Z-(cis) geometric configuration, or a mixture thereof.

10 In the practice of the invention, the amount of the compound incorporated in the pharmaceutical composition may vary widely. Factors considered when determining the precise dose are well known to those skilled in the art. Examples of such factors include, but are not limited to, the subject being treated and 15 the specific pharmaceutical carrier, as well as the route and frequency of administration.

20 A therapeutically effective dose of the compound may comprise about 10 to about 6,000 mg of the active ingredient. In one embodiment, the therapeutically effective dose comprises about 10 to about 3,000 mg. In another embodiment, the therapeutically effective dose comprises about 10 to about 2,000 mg. In a preferred embodiment, the therapeutically effective dose comprises about 10 to about 1,000 mg of the 25 active ingredient. In another embodiment, the therapeutically effective dose comprises about 50 mg to about 500 mg. In a further embodiment, the therapeutically effective dose comprises about 500 to about 4000 mg. In another embodiment, the therapeutically effective dose comprises about 1000 to 30 about 3000 mg. In yet another embodiment, the therapeutically effective dose comprises about 2000 to about 3000 mg. In a preferred embodiment, the therapeutically effective dose

-11-

comprises about 3000 mg.

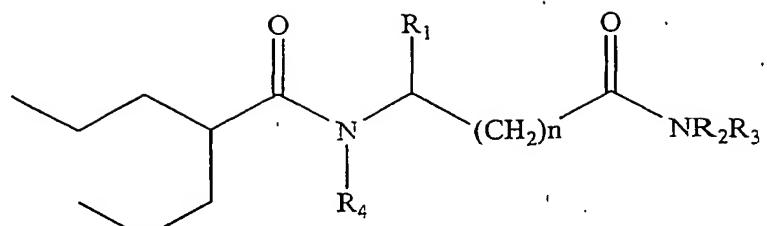
A prophylactically effective dose of the compound may comprise about 10 to about 6,000 mg of the active ingredient. In one embodiment, the prophylactically effective dose comprises about 5 10 to about 3,000 mg. In another embodiment, the prophylactically effective dose comprises about 10 to about 2,000 mg. In a preferred embodiment, the prophylactically effective dose comprises about 10 to about 1,000 mg of the active ingredient. In another embodiment, the prophylactically effective dose comprises about 50 mg to about 500 mg. In a further embodiment, the prophylactically effective dose comprises about 500 to about 4000 mg. In another embodiment, the prophylactically effective dose comprises about 1000 to about 3000 mg. In yet another embodiment, the prophylactically effective dose comprises about 2000 to about 3000 mg. In a preferred embodiment, the prophylactically effective dose comprises about 3000 mg.

The administration of the compound may be effected once daily 20 or up to 6 times per day. Thus, in one embodiment, the administration of the compound may be effected twice a day. In another embodiment, the administration of the compound may be effected 3 times a day. In a further embodiment, the administration of the compound may be effected 4 times a day. 25 In yet another embodiment, the administration of the compound may be effected 5 times a day. In an added embodiment, the administration of the compound may be effected 6 times a day.

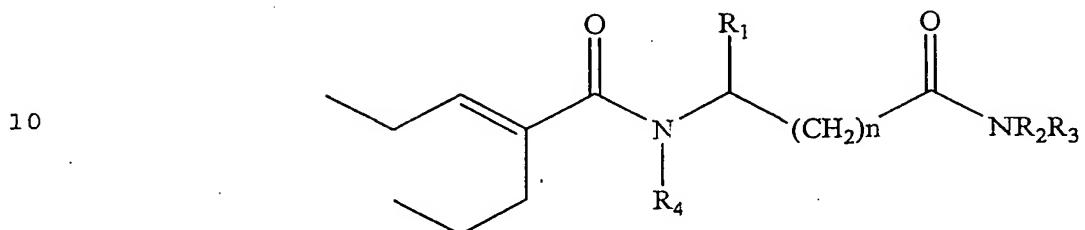
The subject invention also provides a method of treating a 30 subject suffering from pain comprising periodically administering to the subject a therapeutically effective dose of composition comprising a compound having the following

-12-

structure:



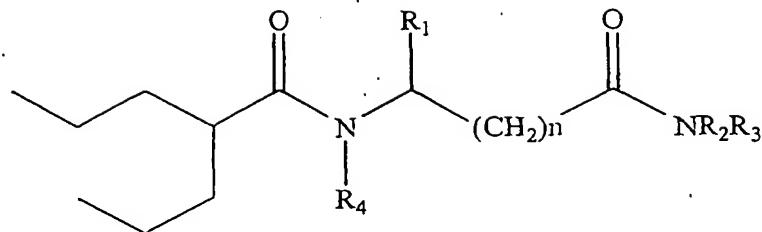
or



wherein R₁, R₂, R₃ and R₄ are independently the same or
 15 different and are hydrogen, a linear or branched C₁-C₆ alkyl group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3, so as to thereby treat the subject's pain.

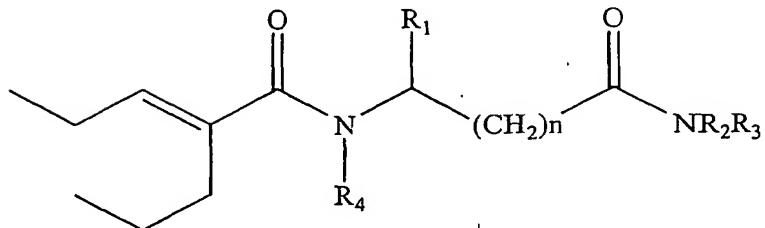
20 Additionally, the subject invention provides a method of preventing pain in a subject predisposed to suffering from pain comprising periodically administering to the subject a prophylactically effective dose of composition comprising a compound having the following structure:

-13-



5

or



10

wherein R₁, R₂, R₃ and R₄ are independently the same or different and are hydrogen, a linear or branched C₁-C₆ alkyl group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3, so as to thereby prevent pain in the subject.

The subject invention also provides compounds containing a valproic acid moiety or a 2-valproenic acid moiety, as well as pharmaceutical compositions comprising these compounds.

The subject invention further provides a pharmaceutical composition comprising the compound and a pharmaceutically acceptable carrier. As used herein, the term "pharmaceutically acceptable carrier" encompasses any of the standard pharmaceutically accepted carriers, such as a phosphate-buffered saline solution, water, suspensions, powders,

-14-

emulsions such as an oil/water emulsion or a triglyceride emulsion, various types of wetting agents, tablets, coated tablets, dissolvable tablets and capsules. An example of an acceptable triglyceride emulsion useful in the intravenous and intraperitoneal administration of the compounds is the 5 triglyceride emulsion commercially known as *Intralipid*®.

Typically, pharmaceutically acceptable carriers contain 10 excipients such as starch, milk, sugar, certain types of clay, gelatin, stearic acid, talc, vegetable fats or oils, gums, glycols, or other known excipients. These carriers may also include flavor and color additives or other ingredients.

In the practice of the invention, the administration of the 15 pharmaceutical composition may be effected by any of the well known methods including, but not limited to, by inhalation, rectal, oral, intravenous, intraperitoneal, parenteral, intramuscular, transdermal, subcutaneous, sublingual, nasal, buccal, pulmonary, vaginal or topical administration.

20 Topical administration can be effected by any method commonly known to those skilled in the art. These methods include, but are not limited to, incorporation of the pharmaceutical composition into creams, gels, ointments, transdermal patches or other topical formulations and delivery systems.

25 When the compound is introduced orally, it may be mixed with other food forms and consumed in solid, semi-solid, suspension or emulsion form. The compound may be administered as sprinkles. In one embodiment, the oral composition may be enterically-coated. Use of enteric coatings are well known in 30 the art. Commonly known enteric coatings include Eudragit S and Eudragit L (Lehman, K., 1971; Lehman, K. 1973; Handbook of

-15-

Pharmaceutical Excipients, 2nd ed.).

The invention encompasses a pharmaceutical composition as hereinabove described wherein the carrier is a solid and the composition is a tablet. The invention also encompasses a pharmaceutical composition as hereinabove described wherein the carrier is a gel and the composition is a suppository. The invention further encompasses a pharmaceutical composition as hereinabove described wherein the carrier is a liquid and the composition is a solution.

10

The following Experimental Details are set forth to aid in an understanding of the invention, and are not intended, and should not be construed, to limit in any way the invention set forth in the claims which follow thereafter.

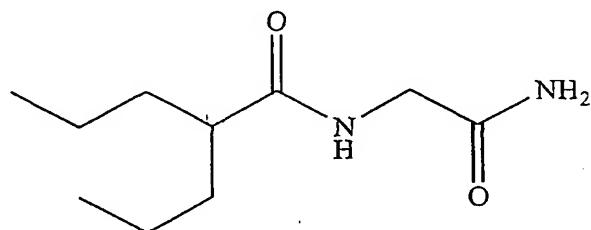
-16-

Experimental Details

I. Synthesis of Compounds

Compound 1 N-(2-n-propylpentanoyl)glycinamide

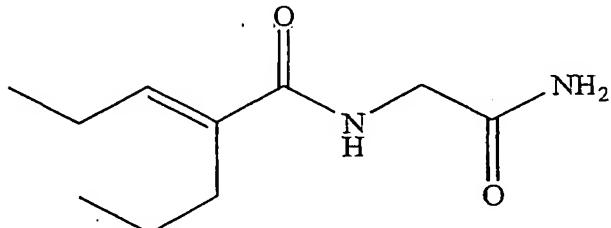
5



10 Compound 1 was prepared as disclosed by U.S. Patent 5,585,358.

Compound 2 N-(2-n-propylpent-2-enoyl)glycinamide

15



Compound 2 was prepared as disclosed by U.S. Patent 5,585,358.

20

II. Experimental Examples

Example 1

25 The anti-pain effects of Compounds 1 and 2 are evaluated in a model for traumatic nerve injury. The specific model is the

-17-

rat constriction injury model, a commonly accepted model for the evaluation of the potential of a compound to treat chronic neuropathic pain. The end point is whether a compound can reverse cold allodynia in rats following a neuropathic injury. MC may be used as the control.

5

RESULTS

Compounds 1 and 2 reverse cold allodynia in the chronic constriction injury model of neuropathic pain with ED_{50} values of less than 500 mg/kg. The effective dose is below that which has been previously found to be the median ataxic dose (also referred to as the minimal neurotoxic dose). MC may be used as the control.

15

DISCUSSION

10

The results indicate that Compounds 1 and 2 are effective for the treatment of pain. Thus, the disclosed valproic acid amides and 2-valproenic acid amides are effective for the treatment or prevention of pain, including neuropathic pain.

Example 2

25

The potential of Compound 1 to serve as an anti-pain agent was studied in the Chung model (Kim, S.H. and Chung, J.M.). This model is known as a reliable model, predictive for human pain. (Kim, S.H. and Chung, J.M.). In this model, spinal nerves L5 and L6 of the rat are tightly ligated and cut in order to induce neuropathic pain. Male Sabra rats weighing 250-275 g

-18-

were used throughout the study. Under xylazine-ketamine anesthesia, both the L5 and L6 spinal nerves of one side of the rat were tightly ligated and cut. Pain behavior was measured following operation in all groups using withdrawal latencies of the hind paw to mechanical stimulation with von Frey filaments (tactile allodynia). The mechanical sensitivity of the foot was quantified by the occurrence of foot withdrawal in response to normally innocuous mechanical stimuli. Eight different von Frey filaments ranging from 0.6 to 26 g were used.

10

The efficacy (antiallodynic effect) of the compound was evaluated (300 mg/kg, *per os* (oral)) in eight rats at days 7 and 14 postoperation in a double-blind randomized crossover manner. The testing included estimation of time of peak effect following drug administration and measurement of the ability of the compound to decrease tactile allodynia.

RESULTS

20 The compound reversed tactile allodynia in the Chung model in comparison to vehicle (MC-methyl cellulose). It was found that the compound reversed the tactile allodynia and therefore, the hind paw withdrawal occurred at higher thresholds. The time of peak effect was 60 minutes. At a statistically significant level, the compound prevented tactile allodynia when compared to vehicle 60 minutes ($p=0.0207$) and 120 minutes ($p=0.0102$) following drug administration (Mann-Whitney test). The results are shown in Figure 1.

30

DISCUSSION

The results demonstrated that Compound 1 is effective for the treatment of pain. Thus, the disclosed valproic acid amides are effective for the treatment or prevention of pain, including neuropathic pain.

Example 3

Evaluation of the anti-headache effects of Compounds 1 and 2 are followed in the migraine model of Moskowitz (Suzzi, M.C. and Moskowitz, M.A.). In this model, neurogenic inflammation results in the leakage of plasma proteins into the dura matter (plasma protein extravasation), which can be quantified by measuring the leakage of radioactive albumin (Suzzi, M.C. and Moskowitz, M.A.).

RESULTS

The results of the experiment employing Compounds 1 and 2 separately are displayed in Table 1. Individually, Compounds 1 and 2 inhibit plasma protein extravasation as compared to the control group (Table 1).

-20-

Table 1: Inhibition of Plasma Protein Extravasation by Compounds 1 and 2

| | Control | Compound 1 | Compound 2 |
|---|---------|------------|------------|
| Percent Extravasation Compared to Control | 100 | < 100 | < 100 |

DISCUSSION

10 The Moskowitz model, which is a well-accepted model of migraines (Suzzi, M.C. and Moskowitz, M.A.), shows that Compounds 1 and 2 inhibit plasma protein extravasation. Thus, the disclosed valproic acid amides and 2-valproenic acid amides are effective for the treatment or prevention of headache 15 disorders, such as migraines.

References

U.S. Patent No. 5,585,358, Bialer et al., issued December 6, 1996.

5 Attal', N. 1999, Mechanism of action and rationale for use of antiepileptic drugs in: International Congress and Symposium Series 241 The Royal Society of Medicine Press Limited Ed. JM Pellock.

10 Handbook of Pharmaceutical Excipients, 2nd ed., A.H. Kibbe, Ph.D., ed., American Pharmaceutical Association and Pharmaceutical Press, Washington, D.C., 2000.

International Headache Society, 1988.

Kim, S.H. and Chung, J.M., 1992, Pain 50: 355-363.

Lehman, K., Acrylic Coatings in Controlled Release Tablet Manufacture, Manufacturing Chemist and Aerosol News, 1973, 39.

15 Lehman, K., Programmed Drug Release from Oral Program Forms, Pharma. Int., vol. ISS 3, 1971, 34-41.

Moskowitz, M.A. and Cutrer, F.M., Sumatriptan: a receptor-targeted treatment for migraines. Ann. Rev. Med., 1993: 44:145-154.

20 Silberstein, S.D. et al., 1998, Headache in Clinical Practice, Pub. Isis Medical Media, Oxford.

Suzzi, M.C. and Moskowitz, M.A., The Antimigraine Drug, Sumetriptan (GR43175), Selectively Blocks Neurogenic Plasma Extravasation from Blood Vessels in Dura Mater, 1990, Br. J. Pharmcol., 99: 202-206.

25 Tremont-Lukats, I. et al., Anticonvulsants for Neuropathic Pain, Drugs, 2000, 60: 1029.

Woolf, C. and Mannion, R., Neuropathic Pain: Aetiology, Symptoms, Mechanisms and Management, Lancet, 1999, 353: 1959.

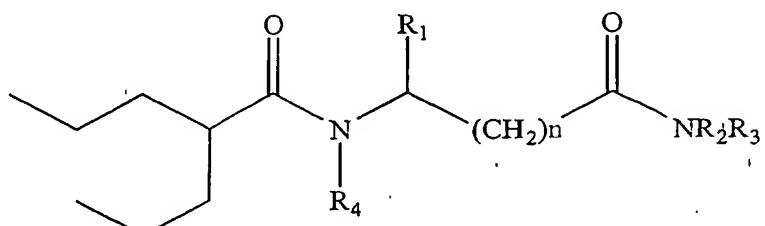
-22-

The Merck Manual of Diagnosis and Therapy, 17th Ed., 1999, M. Beers and R. Berkow, eds., Merck Research Laboratories, Whitehouse Station, N.J.

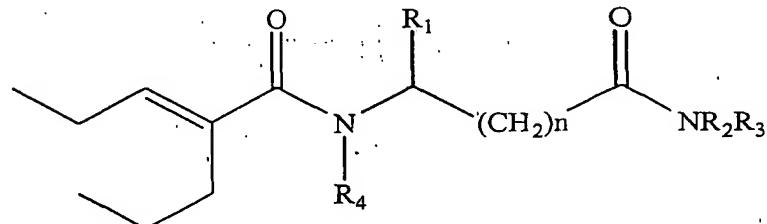
Principles of Neurology, 6th Ed., 1997, Adams, R.D. and Victor, M., McGraw-Hill, Inc., 148-159.

What is claimed

1. A method of treating a subject suffering from pain comprising periodically administering to the subject a therapeutically effective dose of a compound having the following structure:



or



20 wherein R₁, R₂, R₃ and R₄ are independently the same or different and are hydrogen, a linear or branched C₁-C₆ alkyl group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3, so as to thereby treat the subject's pain.

25

2. The method of claim 1, wherein one or more of R₁, R₂, R₃ or R₄ is a linear chain C₁-C₆ alkyl group.

3. The method of claim 1, wherein one or more of R₁, R₂, R₃ or R₄ is a branched chain C₁-C₆ alkyl group.

4. The method of claim 1, wherein one or more of R₁, R₂, R₃ or R₄ is a benzyl, alkylbenzyl, hydroxybenzyl, alkoxy carbonylbenzyl, aryloxy carbonylbenzyl, carboxybenzyl, nitrobenzyl, cyanobenzyl, or halobenzyl group.

10 5. The method of claim 1, wherein one or more of R₁, R₂, R₃ or R₄ is a phenyl, naphthyl, anthracenyl, pyridinyl, indolyl, furanyl, alkylphenyl, hydroxyphenyl, alkoxy carbonylphenyl, aryloxy carbonylphenyl, nitrophenyl, cyanophenyl, halophenyl group, mercaptophenyl, or aminophenyl group.

15 6. The method of claim 1, wherein the pain is acute pain.

7. The method of claim 1, wherein the pain is chronic pain.

20 8. The method of claim 1, wherein the pain is somatogenic pain.

25 9. The method of claim 8, wherein the somatogenic pain is neuropathic pain.

10. The method of claim 1, wherein the subject is a human being.

-25-

11. The method of claim 1, wherein the administration is oral, parenteral, intraperitoneal, intravenous, intramuscular, transdermal, subcutaneous, topical or rectal administration.

5

12. The method of claim 1, wherein the administration is by inhalation, sublingual, nasal, buccal, pulmonary or vaginal administration.

10 13. The method of claim 1, wherein the periodic administration is effected daily.

14. The method of claim 1, wherein the periodic administration is effected less than or equal to six
15 times a day.

15. The method of claim 14, wherein the periodic administration is effected six times a day.

20 16. The method of claim 1, wherein the therapeutically effective dose is an amount from about 10 mg to about 6,000 mg.

25 17. The method of claim 16, wherein the therapeutically effective dose is an amount from about 500 mg to about 4,000 mg.

18. The method of claim 16, wherein the therapeutically

-26-

effective dose is an amount from about 10 mg to about 3,000 mg.

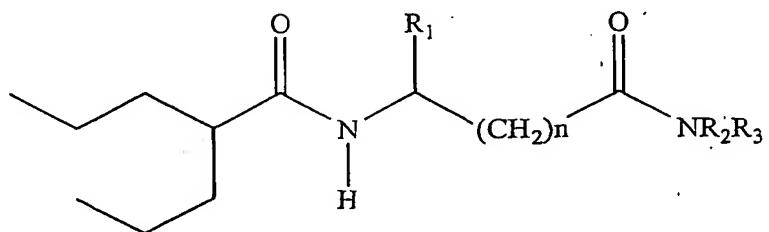
19. The method of claim 18, wherein the therapeutically effective dose is about 3,000 mg.

5

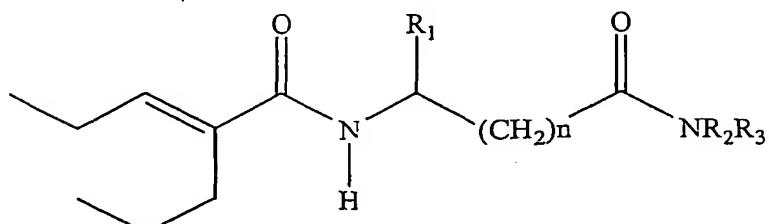
20. The method of claim 18, wherein the therapeutically effective dose is an amount from about 10 mg to about 1,000 mg.

10 21. The method of claim 20, wherein the therapeutically effective dose is an amount from about 50 mg to about 500 mg.

15 22. The method of claim 1, wherein the compound has the following structure:



or



-27-

23. The method of claim 22, wherein the compound is N-(2-n-propylpentanoyl)glycinamide.

24. The method of claim 23, wherein the therapeutically effective dose is 3000 mg/day and the pain is neuropathic pain.

25. The method of claim 22, wherein the compound is N-2(-n-propylpent-2-enoyl)glycinamide.

10

26. The method of claim 22, wherein one or more of R₁, R₂, R₃ or R₄ is a linear chain C₁-C₆ alkyl group.

15

27. The method of claim 22, wherein one or more of R₁, R₂, R₃ or R₄ is a branched chain C₁-C₆ alkyl group.

20

28. The method of claim 22, wherein one or more of R₁, R₂, R₃ or R₄ is aralkyl group is a benzyl, alkylbenzyl, hydroxybenzyl, alkoxy carbonylbenzyl, aryloxycarbonylbenzyl, carboxybenzyl, nitrobenzyl, cyanobenzyl, or halobenzyl group.

25

29. The method of claim 22, wherein one or more of R₁, R₂, R₃ or R₄ is a phenyl, naphthyl, anthracenyl, pyridinyl, indolyl, furanyl, alkylphenyl, hydroxyphenyl, alkoxy carbonylphenyl, aryloxycarbonylphenyl, nitrophenyl, cyanophenyl, halophenyl group, mercaptophenyl, or aminophenyl group.

-28-

30. The method of claim 22, wherein the pain is acute pain.
31. The method of claim 22, wherein the pain is chronic pain.
32. The method of claim 22, wherein the pain is somatogenic pain.
33. The method of claim 32, wherein the somatogenic pain is neuropathic pain.
34. The method of claim 22, wherein the subject is a human being.
35. The method of claim 22, wherein the administration oral, parenteral, intraperitoneal, intravenous, intramuscular, transdermal, subcutaneous, topical or rectal administration.
36. The method of claim 22, wherein the administration is by inhalation, sublingual, nasal, buccal, pulmonary or vaginal administration.
37. The method of claim 22, wherein the periodic administration is effected daily.
38. The method of claim 22, wherein the periodic administration is effected less than or equal to six times a day.

-29-

39. The method of claim 38, wherein the periodic administration is effected six times a day.

40. The method of claim 22, wherein the therapeutically effective dose is an amount from about 10 mg to about 5 6,000 mg.

41. The method of claim 40, wherein the therapeutically effective dose is an amount from about 500 mg to about 4,000 mg.

10

42. The method of claim 40, wherein the therapeutically effective dose is an amount from about 10 mg to about 3,000 mg.

15

43. The method of claim 42, wherein the therapeutically effective dose is about 3,000 mg.

20

44. The method of claim 42, wherein the therapeutically effective dose is an amount from about 10 mg to about 1,000 mg.

45. The method of claim 44, wherein the therapeutically effective dose is an amount from about 50 mg to about 500 mg.

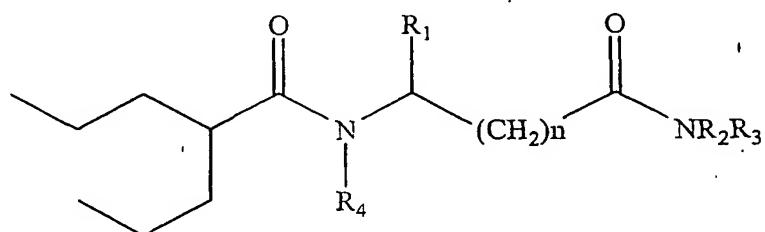
25

46. A method of treating a subject suffering from neuropathic pain comprising administering to the subject 500 mg of N-(2-n-propylpentanoyl)glycinamide six times per day so as to thereby treat the subject's neuropathic pain.

-30-

47. A method of preventing pain in a subject predisposed to suffering from pain comprising periodically administering to the subject a prophylactically effective dose of a compound having the following structure:

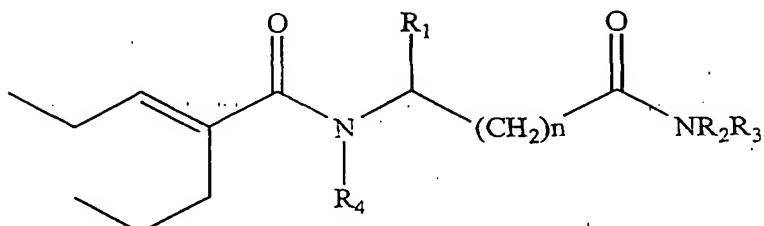
5



10

or

15



wherein R₁, R₂, R₃ and R₄ are independently the same or different and are hydrogen, a linear or branched C₁-C₆ alkyl group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3, so as to thereby prevent pain in the subject.

25 48. The method of claim 47, wherein one or more of R₁, R₂, R₃ or R₄ is a linear chain C₁-C₆ alkyl group.

-31-

49. The method of claim 47, wherein one or more of R₁, R₂, R₃ or R₄ is a branched chain C₁-C₆ alkyl group.

50. The method of claim 47, wherein one or more of R₁, R₂, R₃ or R₄ is a benzyl, alkylbenzyl, hydroxybenzyl, alkoxy carbonylbenzyl, aryloxy carbonylbenzyl, carboxybenzyl, nitrobenzyl, cyanobenzyl, or halobenzyl group.

51. The method of claim 47, wherein one or more of R₁, R₂, R₃ or R₄ is a phenyl, naphthyl, anthracenyl, pyridinyl, indolyl, furanyl, alkylphenyl, hydroxyphenyl, alkoxy carbonylphenyl, aryloxy carbonylphenyl, nitrophenyl, cyanophenyl, halophenyl group, mercaptophenyl, or aminophenyl group.

15

52. The method of claim 47, wherein the pain is acute pain.

53. The method of claim 47, wherein the pain is chronic pain.

20

54. The method of claim 47, wherein the pain is somatogenic pain.

55. The method of claim 54, wherein the somatogenic pain is neuropathic pain.

25

56. The method of claim 47, wherein the subject is a human being.

-32-

57. The method of claim 47, wherein the administration is oral, parenteral, intraperitoneal, intravenous, intramuscular, transdermal, subcutaneous, topical or rectal administration.

5 58. The method of claim 47, wherein the administration is by inhalation, sublingual, nasal, buccal, pulmonary or vaginal administration.

10 59. The method of claim 47, wherein the periodic administration is effected daily.

60. The method of claim 47, wherein the periodic administration is effected less than or equal to six times a day.

15 61. The method of claim 60, wherein the periodic administration is effected six times a day.

20 62. The method of claim 47, wherein the prophylactically effective dose is an amount from about 10 mg to about 6,000 mg.

25 63. The method of claim 62, wherein the prophylactically effective dose is an amount from about 500 mg to about 4,000 mg.

64. The method of claim 62, wherein the prophylactically effective dose is an amount from about 10 mg to about 3,000 mg.

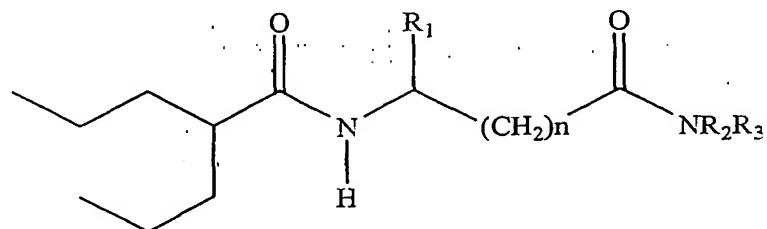
-33-

65. The method of claim 64, wherein the prophylactically effective dose is about 3,000 mg.

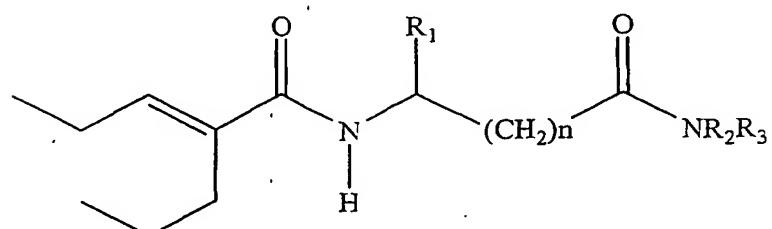
5 66. The method of claim 64, wherein the prophylactically effective dose is an amount from about 10 mg to about 1,000 mg.

10 67. The method of claim 66, wherein the prophylactically effective dose is an amount from about 50 mg to about 500 mg.

68. The method of claim 47, wherein the compound has the following structure:



or



25

-34-

69. The method of claim 68, wherein the compound is N-(2-n-propylpentanoyl)glycinamide.

70. The method of claim 69, wherein the prophylactically effective dose is 3000 mg/day and the pain is neuropathic
5 pain.

71. The method of claim 68, wherein the compound is N-2(-n-propylpent-2-enoyl)glycinamide.

10 72. The method of claim 68, wherein one or more of R₁, R₂, R₃ or R₄ is a linear chain C₁-C₆ alkyl group.

73. The method of claim 68, wherein one or more of R₁, R₂, R₃ or R₄ is a branched chain C₁-C₆ alkyl group.

15 74. The method of claim 68, wherein one or more of R₁, R₂, R₃ or R₄ is aralkyl group is a benzyl, alkylbenzyl, hydroxybenzyl, alkoxycarbonylbenzyl, aryloxycarbonylbenzyl, carboxybenzyl, nitrobenzyl, cyanobenzyl, or halobenzyl group.
20

75. The method of claim 68, wherein one or more of R₁, R₂, R₃ or R₄ is a phenyl, naphthyl, anthracenyl, pyridinyl, indolyl, furanyl, alkylphenyl, hydroxyphenyl, alkoxycarbonylphenyl, aryloxycarbonylphenyl, nitrophenyl, cyanophenyl, halophenyl group, mercaptophenyl, or aminophenyl group.
25

76. The method of claim 68, wherein the pain is acute pain.

-35-

77. The method of claim 68, wherein the pain is chronic pain.

78. The method of claim 68, wherein the pain is somatogenic pain.

5

79. The method of claim 78, wherein the somatogenic pain is neuropathic pain.

80. The method of claim 68, wherein the subject is a human being.

10

81. The method of claim 68, wherein the administration oral, parenteral, intraperitoneal, intravenous, intramuscular, transdermal, subcutaneous, topical or rectal administration.

15

82. The method of claim 68, wherein the administration is by inhalation, sublingual, nasal, buccal, pulmonary or vaginal administration.

20

83. The method of claim 68, wherein the periodic administration is effected daily.

25

84. The method of claim 68, wherein the periodic administration is effected less than or equal to six times a day.

85. The method of claim 68, wherein the periodic

-36-

administration is effected six times a day.

86. The method of claim 68, wherein the prophylactically effective dose is an amount from about 10 mg to about 6,000 mg.

5

87. The method of claim 86, wherein the prophylactically effective dose is an amount from about 500 mg to about 4,000 mg.

10

88. The method of claim 86, wherein the prophylactically effective dose is an amount from about 10 mg to about 3,000 mg.

15

89. The method of claim 88, wherein the prophylactically effective dose is about 3,000 mg.

90. The method of claim 88, wherein the prophylactically effective dose is an amount from about 10 mg to about 1,000 mg.

20

91. The method of claim 90, wherein the prophylactically effective dose is an amount from about 50 mg to about 500 mg.

25

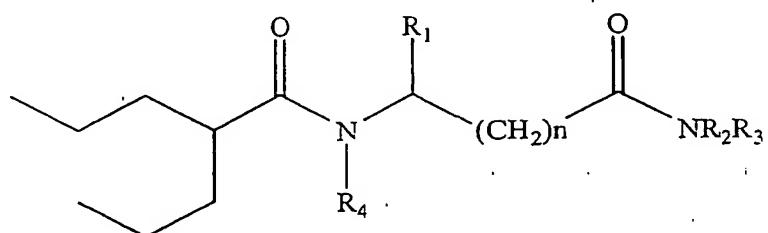
92. A method of preventing neuropathic pain in a subject predisposed to suffering from neuropathic pain comprising administering to the subject 500 mg of N-(2-n-propylpentanoyl)glycinamide six times per day so as to thereby prevent the neuropathic pain in the subject.

-37-

93. A method of treating a subject suffering from pain comprising periodically administering to the subject a pharmaceutical composition comprising a therapeutically effective dose a compound having the following structure:

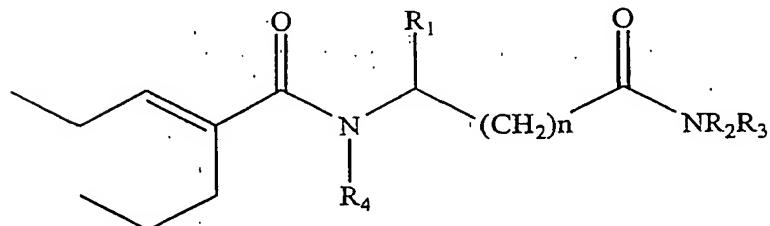
5

10



or

15



20

wherein R₁, R₂, R₃ and R₄ are independently the same or different and are hydrogen, a linear or branched C₁-C₆ alkyl group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3,

25

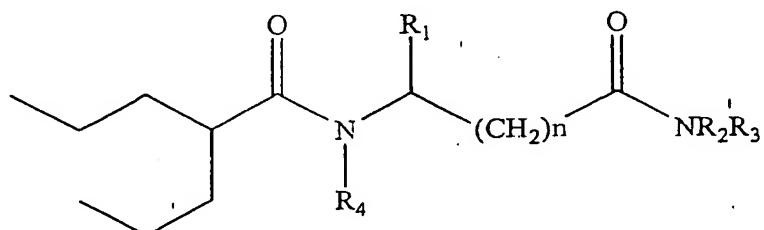
and a pharmaceutically acceptable carrier, so as to thereby treat the subject's pain.

94. A method of preventing pain in a subject predisposed to

-38-

suffering from pain comprising periodically administering to the subject a composition comprising a prophylactically effective dose of a compound having the following structure:

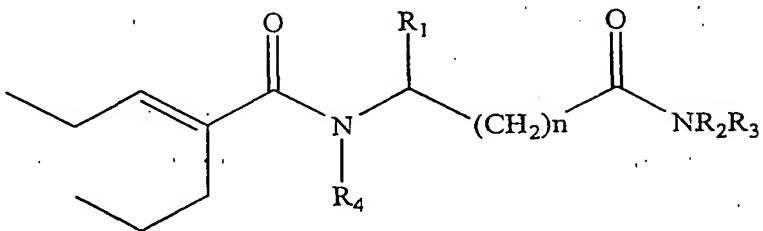
5



10

or

15



20

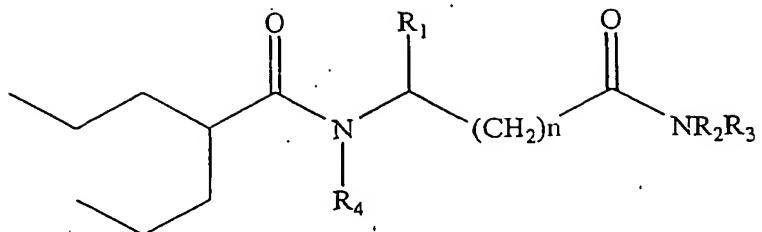
wherein R₁, R₂, R₃ and R₄ are independently the same or different and are hydrogen, a linear or branched C₁-C₆ alkyl group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3,

and a pharmaceutically acceptable carrier, so as to thereby prevent pain in the subject.

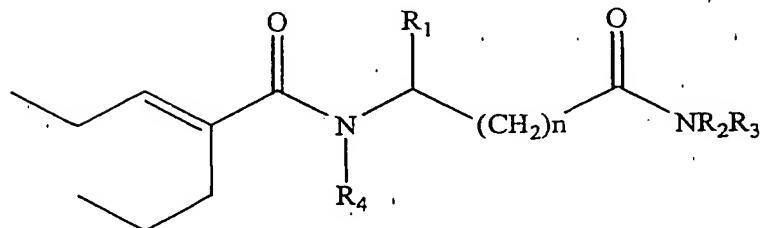
25

95. A method of treating a subject suffering from a headache disorder comprising periodically administering to the subject a therapeutically effective dose of a compound having the following structure:

-39-



or

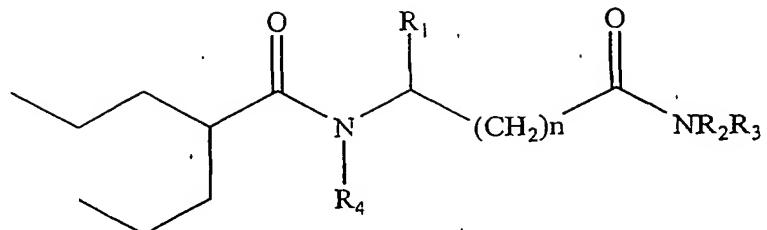


wherein R₁, R₂, R₃ and R₄ are independently the same or different and are hydrogen, a linear or branched C₁-C₆ alkyl group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3, so as to thereby treat the headache disorder.

15

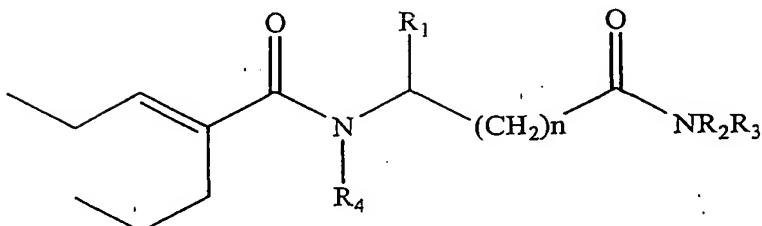
20 96. A method of preventing a headache disorder in a subject predisposed to suffering from a headache disorder comprising periodically administering to the subject a prophylactically effective dose of a compound having the following structure:

-40-



5

or



10

wherein R₁, R₂, R₃ and R₄ are independently the same or different and are hydrogen, a linear or branched C₁-C₆ alkyl group, an aralkyl group, or an aryl group, and n is an integer which is greater than or equal to 0 and less than or equal to 3, so as to thereby prevent the headache disorder in the subject.

15

